# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Food and Drug Administration

#### [Docket No. FDA-2016-N-0001]

### Request for Nominations on the Technical Electronic Product Radiation Safety Standards Committee

**AGENCY:** Food and Drug Administration, HHS.

#### ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is requesting any industry organizations interested in participating in the selection of voting industry representatives to serve on the Technical Électronic Product Radiation Safety Standards Committee for the Center for Devices and Radiological Health to notify FDA in writing. FDA is also requesting nominations for voting industry representatives to serve on the Technical Électronic Product Radiation Safety Standards Committee. A nominee may either be self-nominated or nominated by an organization to serve as a voting industry representative. Nominations will be accepted for current vacancies effective with this notice.

**DATES:** Any industry organization interested in participating in the selection of an appropriate voting member to represent industry interests must send a letter stating that interest to FDA by *February 18, 2016*, (see sections I and II of this document for further details). Concurrently, nomination materials for prospective candidates should be sent to FDA by February 18, 2016.

**ADDRESSES:** All statements of interest from industry organizations interested in participating in the selection process of voting industry representative nomination should be sent to Margaret Ames (see FOR FURTHER INFORMATION **CONTACT**). All nominations for voting industry representatives may be submitted electronically by accessing the FDA Advisory Committee Membership Nomination Portal: https:// www.accessdata.fda.gov/scripts/ FACTRSPortal/FACTRS/index.cfm or by mail to Advisory Committee Oversight and Management Staff, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 5103, Silver Spring, MD 20993-0002. Information about becoming a member of an FDA advisory committee can also be obtained by visiting FDA's Web site at http:// www.fda.gov/AdvisoryCommittees/ default.htm.

FOR FURTHER INFORMATION CONTACT: Margaret Ames, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, rm. 5234, Silver Spring, MD 20993, 301–796–5960, FAX: 301– 847–8505, margaret.ames@fda.hhs.gov.

**SUPPLEMENTARY INFORMATION:** The Agency requests nominations for voting industry representatives to the following advisory committee:

#### I. Technical Electronic Product Radiation Safety Standards Committee

This Committee provides advice and consultation to the Commissioner of Food and Drugs on the technical feasibility, reasonableness, and practicability of performance standards for electronic products, to control the emission of radiation from such products, and may recommend electronic product radiation safety standards to the Commissioner for consideration.

#### **II. Selection Procedure**

Any industry organization interested in participating in the selection of an appropriate voting member to represent industry interests should send a letter stating that interest to the FDA contact (see FOR FURTHER INFORMATION CONTACT) within 30 days of publication of this document (see DATES). Within the subsequent 30 days, FDA will send a letter to each organization that has expressed an interest, attaching a complete list of all such organizations, and a list of all nominees along with their current resumes. The letter will also state that it is the responsibility of the interested organizations to confer with one another and to select a candidate, within 60 days after the receipt of the FDA letter, to serve as the voting member to represent industry interests for the committee. The interested organizations are not bound by the list of nominees in selecting a candidate. However, if no individual is selected within 60 days, the Commissioner will select the voting member to represent industry interests.

### **III. Application Procedure**

Individuals may self-nominate and/or an organization may nominate one or more individuals to serve as a voting industry representative. Contact information, a current curriculum vitae, and the name of the committee of interest should be sent to the FDA Advisory Committee Membership Nomination Portal (see **ADDRESSES**) within 30 days of publication of this document (see **DATES**). FDA will forward all nominations to the organizations expressing interest in participating in the selection process for the committee. Persons who nominate themselves as voting industry representatives will not participate in the selection process.

FDA seeks to include the views of women, men, members of all racial and ethnic groups, and individuals with and without disabilities on its advisory committees and, therefore encourages nominations of appropriately qualified candidates from these groups.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to advisory committees.

Dated: January 11, 2016.

### Jill Hartzler Warner,

Associate Commissioner for Special Medical Programs.

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# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Food and Drug Administration

[Docket No. FDA-2015-N-0001]

### Circulatory System Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

**AGENCY:** Food and Drug Administration, HHS.

### ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

*Name of Committee:* Circulatory System Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on

FDA's regulatory issues. Date and Time: The meeting will be

held on March 15 and 16, 2016, from 8 a.m. to 6 p.m.

*Location:* Holiday Inn Gaithersburg, Ballroom, Two Montgomery Village Ave., Gaithersburg, MD 20879. The hotel telephone number is 301–948– 8900.

Contact Person: Dimitrus Culbreath, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, rm. 3610, Silver Spring, MD 20993– 0002, 301–796–6872, Dimitrue Culbreath of da has gov, en

*Dimitrus.Culbreath@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously